

K240040 VistaPano S Ceph 2.0 (VistaPano S Ceph)Oct 25, 2024
294 days to decisionK240040 · Product code: **MUH** · Radiology
Source: <https://www.510kdatabase.net/k240040/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Extraoral Source, Digital (MUH)
Date received	Jan 5, 2024
Decision date	Oct 25, 2024
Days to decision	294 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary
Other names	VistaPano S 2.0 (VistaPano S); ProVecta S-Pan Ceph 2.0 (ProVecta S-Pan Ceph); ProVecta S-Pan 2.0 (ProVecta S-Pan)

APPLICANT

Company	VATECH Co., Ltd.
Location	Sound Beach, NY, US
Contact	Daniel Kim
510(k) history	39 submissions · 39 cleared · 2008-2025

REGULATORY CONSULTANT

Consulting firm	Mtech Group, LLC
Contact	Dave Kim

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)**510k Database** - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k240040/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 13, 2026